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2	Oil biodegradation in permeable marine sediments: Effects of benthic pore-water
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### 38 Abstract

Oil spills have been recognized as among the worst kinds of environmental disasters, causing 39 severe coastal ecological and economic damages. Although benthic flow and solute fluxes are 40 known to have strong impacts on fate and transport of oil deposited within marine sediments, 41 42 their endogenous mechanisms still remain to be uncovered. In this paper, simulations of flow and solute transport processes along with hydrocarbon biodegradation were conducted in a 43 cylindrical benthic chamber system to investigate influences of benthic hydrodynamics on oil 44 45 biodegradation in permeable marine sediments. Results show that ripple-flow interactions create subsurface recirculation cells whereby seawater infiltrates into the benthic sediments at 46 47 ripple troughs while groundwater discharges near the crests. It results in a spatially varied oil 48 biodegradation rate in marine sediments. Significant oil biodegradation occurs near sediment ripple troughs due to direct oxygen recharge, while biodegradation of oil deposited uphill 49 50 becomes slow due to limited oxygen replenishment. Oil biodegradation decreases subsurface oxygen content, and consequently impedes discharge of oxygen from benthic sediments. Our 51 52 results reveal a dynamic interaction between oil biodegradation and benthic flow and solute transport processes, which has strong implications for predicting oil persistence and 53 biodegradation within marine sediments and its associated impacts on benthic 54 55 biogeochemical processes.

### 57 **1. Introduction**

Oil spills pose severe threats to marine coastal areas worldwide, deteriorating shoreline 58 59 environments and causing severe ecological and economic damage. The 1989 Exxon Valdez oil spill contaminated around 800 km of shorelines within Prince William Sound, Alaska 60 [Michel and Hayes, 1999; Short et al., 2004; Boufadel et al., 2016]. In 2002, the Prestige oil 61 62 spill occurred off the coast of Galicia, Spain, polluting over 600 sandy beaches along the Spanish Atlantic coast [Fernández-Fernández et al., 2011; Acosta-González et al., 2015]. The 63 64 2010 Deepwater Horizon oil spill led to the release of an estimated 3.19 million barrels of crude oil into the Gulf of Mexico and an estimated 22,000 tons of weathered oil washed up 65 on northern Gulf of Mexico beaches [Michel et al., 2013; Boufadel et al., 2014; Geng et al., 66 67 2021a]. Incidents of these magnitudes resulted in serious consequences on marine ecosystems, degrading abundance and diversity of benthic communities [Glemarec and Hussenot, 1981; 68 Teal and Howarth, 1984]. The toxicity in oils is primarily from aromatic hydrocarbons, 69 particularly polycyclic aromatic hydrocarbons (PAHs) [Neff, 2002]. Marine organisms are 70 found to be affected by PAHs at every life stage, from free floating embryos and larvae to 71 72 sexually mature adults [Adzigbli and Yuewen, 2018], and across species types. For instance, 73 crustaceans can be exposed to oil through digging into oiled sediments, ingestion with food, and direct interaction [Cormack et al., 2011; Montagna et al., 2013], and often experience 7475 high mortality after oil spills [McCay et al., 2001]. Bodkin et al. [2012] reported consequent impacts of the Exxon Valdez oil spill on the decrease in sea otter subpopulation rate in the 76 heavily oiled northern Knight Island area of Prince William Sound. Reuscher et al. [2017] 77 investigated impacts of the Deepwater Horizon oil spill on benthic infauna of the northern 7879 Gulf of Mexico continental slope. Their study found that oil originating from the Deepwater Horizon reached the seafloor and had a persistent negative impact on diversity of soft-bottom, 80 deep-sea benthic communities. While signs of recovery for some benthic community 81

variables are shown, full recovery had not yet occurred even four years after the spill.

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There are various driving mechanisms that can cause sedimentation of spilled oil onto 84 85 nearshore and benthic sediments [Teal and Howarth, 1984; Boufadel et al., 2019]. Mixing of spilled oil with sediments often occurs in shallow water by breaking waves and further 86 87 transport to deeper water is driven by subsequent currents (Figure 1A) [Owens and Lee, 2003; 88 O'Laughlin et al., 2017]. Oil could also attach onto particulate matter suspended in the water column and subsequently sink in deeper water [Gong et al., 2014; Shan et al., 2020]. Pore-89 90 water advection within marine sediments is often generated by the pressure gradients at the 91 sediment surface that occur when bottom current flow is deflected by topographical structures 92 such as mounds or ripples [Huettel and Gust, 1992; Janssen et al., 2005a]. It has been proven 93 to play a critical role in biogeochemical processes in marine benthic sediments, including 94 carbon mineralization, nutrient recycling, and recycled primary production [Janssen et al., 95 2005a; Geng et al., 2021b]. Tracer studies illustrated that ripple-flow interactions generate 96 groundwater recirculation cells in marine sediments with flows infiltrating in the ripple troughs and exfiltrating from crests (Figure 1B) [Webb and Theodor, 1968; Shum, 1992]. 97 98 Such seawater-groundwater recirculation brings organic matter and oxygen into the sediment, creates horizontal concentration gradients that can be as strong as the vertical gradients, and 99 100 provides an important mechanism for enhancing mineralization of organic matter in marine 101 sediments [Huettel et al., 1998]. Although advective flow and solute fluxes across the watersediment interface could have strong impacts on fate and transport of oil deposited within 102 103 marine sediments, endogenous mechanisms of benthic flow and solute exchange interacting 104 with oil biodegradation in marine sediments have not been investigated yet.

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106 Investigation of benthic flow and solute exchange has been widely performed with benthic

107 chambers. Experiments involve benthic stirred chambers where the centrifugal-force-induced 108 pressure gradients at the sediment surface generate pore-water advection that circulates the overlying water through the sediment [Tengberg et al., 1995; Glud et al., 1996], and benthic 109 110 landers deployed to the sea floor for direct in-situ measurements and observations [Black et 111 al., 2001; D'Onghia et al., 2018]. Mathematical modeling has also been widely used to investigate flow and solute transport processes within benthic sediments. It allows us to 112 interpret measurement data collected from experiments and examine complex field-like 113 conditions which are difficult to reproduce with laboratory experiments. *Khalili et al.* [1997] 114 115 and [1999] derived a numerical solution of the Darcy-Brinkmann-Lapwood equation to characterize the axisymmetric advective flow in a cylindrical chamber system. Calculations 116 were performed to quantify the flow exchange between the permeable sediment and the 117 118 overlying water. Meysman et al. [2007] conducted numerical simulations to investigate 119 biologically and physically induced flow and tracer dynamics in marine benthic sediments. In their study, the finite element package COMSOL Multiphysics<sup>TM</sup> 3.2a was used to develop 120 121 flow and reactive transport models for characterizations of advective flow and associated fluxes across the sediment-water interface. While considerable efforts have been made on 122 modeling benthic flow and solute exchange processes, interactions between coastal benthic 123 exchange dynamics and oil biodegradation have not been numerically conducted within 124 125 benthic oil-contaminated sediments.

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127 The objective of this paper is to investigate interactions between benthic oil biodegradation 128 and pore-water advection along with solute exchange in marine sediments. We developed a 129 two-dimensional (2D) flow and reactive transport model in cylindrical coordinates for the 130 investigation. The development was based on transformation of our prior developed 131 BIOMARUN model from Cartesian to cylindrical coordinates. We conducted simulations of

groundwater flow, reactive transport of oxygen and a nutrient (nitrogen), and hydrocarbon biodegradation in a cylindrical benthic sediment chamber to characterize the patterns of microbial degradation of oil subjected to benthic flow and solute flux exchange across the water-sediment interface. Particle tracking was further performed to analyze advective flow paths and associated travel time in this chamber system. The results highlight the important role of benthic hydrodynamics in hydrocarbon biodegradation within permeable marine sediments.

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## 140 **2. Methods**

### 141 2.1 BIOMARUN model in cylindrical coordinates

142 The BIOMARUN model in Cartesian coordinates has been developed and validated in our prior studies [Geng et al., 2015; Geng et al., 2016; Geng et al., 2017a]. The model couples 143 the MARUN model [Boufadel et al., 1999; Geng and Boufadel, 2017; Geng et al., 2020], a 2-144D finite element model considering density-dependent flow and solute transport in variably 145 146 saturated media, with the BIOB model, a multiplicative Monod model for hydrocarbon 147biodegradation [Geng et al., 2013; Geng et al., 2014a]. In this paper, we transformed the BIOMARUN model from Cartesian coordinates to cylindrical coordinates, named CYL-148 149 BIOMARUN, to simulate benthic flow and solute transport processes in a cylindrical 150 chamber system. The equation for the conservation of water is written as:

$$\frac{\partial(\beta\phi S)}{\partial t} = \frac{1}{r} \frac{\partial(r\beta K_r \frac{\partial\psi}{\partial r})}{\partial r} + \frac{\partial(\beta K_z \frac{\partial\psi}{\partial z})}{\partial z} + \frac{\partial(\beta^2 K_z)}{\partial z}, \qquad (1)$$

151 where  $\beta$  is the density ratio [-] defined as the ratio of salt-dependent water density to 152 freshwater density [ML<sup>-3</sup>];  $\phi$  is porosity [-], *S* is soil moisture ratio [-],  $\Psi$  is pressure head 153 [L], and  $K_r$  and  $K_z$  are radial and vertical freshwater hydraulic conductivities. 155 The conservative solute (e.g., salt) transport equation can be expressed as:

$$\phi S \frac{\partial c_0}{\partial t} = \beta \phi S \frac{(D_{rr} \frac{\partial c_0}{\partial r} + D_{rz} \frac{\partial c_0}{\partial z})}{r} + \beta \nabla \cdot (\phi S \vec{D} \cdot \nabla c_0) - \vec{q} \cdot \nabla c_0, \qquad (2)$$

where  $c_0$  is concentration, and  $\vec{q}$  is Darcy flux vector given by:

$$\vec{q} = (q_r, q_z) = -(K_r \frac{\partial \psi}{\partial r}, K_z \frac{\partial \psi}{\partial z} + \beta), \qquad (3)$$

157 The term  $\vec{D}$  represents physical dispersion tensor written as

$$\vec{D} = \frac{1}{\|\vec{q}\|} \begin{pmatrix} \alpha_L q_r^2 + \alpha_T q_z^2 & (\alpha_L - \alpha_T) q_r q_z \\ (\alpha_L - \alpha_T) q_r q_z & \alpha_T q_r^2 + \alpha_L q_z^2 \end{pmatrix},$$
(4)

158 where  $\|\vec{q}\| = \sqrt{q_r^2 + q_z^2}$ ,  $\alpha_L$  and  $\alpha_T$  are longitudinal and transverse dispersivities (L), 159 respectively.

160 The reactive solute transport equation can be expressed as:

$$\phi SR_i \frac{\partial c_i}{\partial t} = \beta \phi S \frac{(D_{rr} \frac{\partial c_i}{\partial r} + D_{rz} \frac{\partial c_i}{\partial z})}{r} + \beta \nabla \cdot (\phi S \vec{D} \cdot \nabla c_i) - \vec{q} \cdot \nabla c_i + \phi DR_i r_i,$$
(5)

where  $R_i$  is the retardation coefficient of  $i_{th}$  solute  $c_i$ ;  $r_i$  is a reaction rate representing 161 generation or consumption. Six reactive components of two types of oil components (alkanes 162 163 and polycyclic aromatic hydrocarbons (PAHs)), their corresponding microbial degraders, dissolved oxygen and a nutrient (nitrogen) are considered. The reaction network of these six 164 components is consistent with the BIOB model, where in general microbes consume 165 dissolved oxygen and nutrients for degrading oil compounds following modified Monod 166 kinetics. For example, the growth coefficient of microbial degraders,  $\mu_i$ , can be expressed as 167 follows: 168

$$\mu_{i} = \mu_{\max} \left(1 - \frac{X_{i}}{X_{\max}}\right) \left(\frac{S_{i}}{K_{s} + S_{i}}\right) \left(\frac{N}{K_{N} + N}\right) \left(\frac{O^{4}}{K_{O} + O^{4}}\right), \tag{6}$$

Where i = 1 and 2, denoting alkane and PAH degraders, respectively, and  $\mu_{\text{max}}$  represents maximum growth coefficients (day<sup>-1</sup>). The parameters  $S_i$ , N, and O represent concentrations of substrate, nutrient, and oxygen. The parameters  $K_S$  and  $K_N$  represent the half-saturation concentrations for substrate and nitrogen, respectively (see *Geng et al.* [2014a] and [2015] for details). The CYL-BIOMARUN model was validated by reproducing the experimental results measured in *Khalili et al.* [1999] (Figure 2).

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### 176 2.2 Numerical simulations of oil biodegradation in a benthic chamber system

177The simulated domain was constructed in a cylindrical coordinate system with the radius length of 50 cm and height of 100 cm, representing a cylindrical benthic chamber system. A 178 mesh of 2278 nodes (34 nodes in the radial and 67 nodes in the vertical) was used, resulting 179 in ~1.5 cm spatial resolution. The mesh was made fine enough to meet the criterion for the 180 grid Peclet number to be less than or equal to 2.0 [Zheng and Bennett, 2002]. For the flow 181 simulation, a steady-state radial-symmetric pressure distribution was applied at the top 182 183 boundary of the domain (Figure 3A). This boundary condition is commonly used for 184 simulating the chamber system where the sediment-water interface is flat, and a radialsymmetric stirring pattern establishes in the overlaying water [Glud et al., 1996]. For fate and 185 transport simulations, a Dirichlet boundary condition was used at the top boundary, and a 186 187 Neumann boundary condition with zero dispersive flux was assigned to the rest of the boundaries. The initial concentration of dissolved oxygen and nutrients was assumed to be 188 6.0 mg/L and 1.0 mg/L in sediments, and 1.2 mg/L and 0.2 mg/L in seawater, respectively. 189 To model microbial degradation of oil, the oil composition can be modeled as pseudo-190 191 components with distinct mass fractions and biodegradation rate [Thrift-Viveros et al., 2015].

For simplification, the oil was assumed to be composed of two major groups, long-chain alkanes and polycyclic aromatic hydrocarbons (PAHs), deposited within the top 5 cm of sediments. The concentrations of alkanes and PAHs were assigned as 4.0 mg/kg of sediment and 2.0 mg/kg of sediment, respectively (Figure 3B).

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197 For all the simulations, the numerical model was first run for approximately 100 days without oil deposition until the hydraulic and hydrodynamic regime reached a steady state. The 198 pressure and solute distributions were then used as initial conditions for the simulation with 199 200 oil contamination near the sediment surface, which ran over a simulation time of 300 days. The time step was selected at 10.0 s, resulting in a grid Courant number less than 0.9 (a value 201 202 less than 1.0 is required). The parameter values adopted in the simulations are reported in 203 Table 1. Backward particle tracking was performed to reveal the flow pathways and 204 associated travel time. A particle tracking code, named NEMO-3D, based on the random walk particle tracking algorithm was used [Geng et al., 2014b; Geng et al., 2017b; Geng et 205 206 al., 2020]. The neutrally buoyant particles were released at a 5-cm interval along the sediment surface, driven by the flow fields obtained from the CYL-BIOMARUN simulation. 207

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# **3. Results**

# 210 3.1 Comparison with the experiment data of Khalili et al. [1999]

*Khalili et al.* [1999] conducted experiments to investigate recirculating flow through benthic fluid-sediment interfaces. The experimental set-up is a stationary cylinder that contains a fluid-saturated porous medium with supernatant water on top. The recirculating flow was generated by deploying a rotating lid in the upper fluid region, and further characterized by monitoring dye washout from the sediment. Dye transport was visualized using positron emission tomography (PET), a non-invasive technique which allows to see through an 217 opaque solid matrix, and to obtain full 3-D pictures of washing out dye from the sediment. In their study, digital photographic visualization and associated image processing were also 218 performed to further quantify the near-wall dye-washout depth. Figure 2A shows a 219 220 comparison of normalized radioactivity strength measured at different times by PET along 221 with numerical results obtained by CYL-BIOMARUN model and the model developed in 222 Basu and Khalili [1999]. As shown in the figure, the CYL-BIOMARUN model reasonably 223 captures the dye transport within the sediment column. The results of washout depth show that both numerical models underestimate the first half-hour washout of dye from the 224 225 sediment (Figure 2B). This is most likely due to the loss of some dye from the sediment to the fluid layer when the experiments were initialized, which was not considered in the 226 227 numerical simulations.

228

# 3.2 Simulations of pore-water advection, solute exchange, and oil biodegradation in a benthic chamber system

Surface pressure gradients have a strong influence on flow and transport processes within 231 232 benthic sediments. Figure 3A shows that the pressure gradients imposed along the sediment surface alter subsurface pressure distribution, creating a high-pressure zone at the outer ring 233 234 of the chamber and relatively low-pressure zone in the proximity of the chamber center. The 235 resulting pressure differences transmit through the subsurface and generate pressure gradients 236in both radial and vertical directions. The penetration depth is approximately 20 cm for the simulations considered in this paper. Figures 3C and 3D show that an upper plume is 237 238 generated in the subsurface with seawater-derived chemical properties that are distinguished from ambient groundwater. This is expected as the formation of the plume is due to seawater 239 infiltration. The plume has a deeper expansion beneath the high-pressure zone, and becomes 240 shallower as it moves to the relatively low-pressure zone. The concentration of oxygen 241

gradually decreases from 6.0 mg/L to 1.0 mg/L and concentration of nutrients gradually increases from 0.2 mg/L to 1.2 mg/L in the proximity of the plume edge, indicating substantial mixing between seawater and groundwater in benthic sediments. The steady-state simulations show the maximum washout depth by seawater is ~20 cm while the thickness of the seawater-groundwater mixing zone is ~25 cm.

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248Particle tracking results show the pressure gradients create a recirculation cell whereby seawater infiltrates into the benthic sediments through the high-pressure zone and 249 250 groundwater discharge occurs along the low-pressure zone (Figure 4). Figure 4A shows the advective flow pathways within the recirculation cell simulated by a backward particle 251 tracking scheme. Infiltration flow occurs near the outer ring of the chamber between R = 40252 253cm and 50 cm, while exfiltration flow is present at the inner part of the chamber between R =0 and 40 cm. The particles migrating along deeper flow paths experience longer travel time 254 (Figure 4B). In particular, the travel time of the particles discharged near the location of R =2552562 cm is about 100 years. The particles' travel time drops exponentially as their discharge 257 locations get close to the infiltration zone due to shorter flow paths.

258

Oil biodegradation demonstrates a large spatial variation in benthic sediments. The 259260 concentration contour of alkanes shows that two biodegradation hotspots appear in the oiled 261 sediment layer after 30 days near the top within the infiltration zone and near the bottom within the exfiltration zone, respectively (Figure 5A). The spatial extent of these two hotspots 262 gradually expands with time (e.g., time t = 60 days and 100 days). After 300 days, nearly all 263 264 the alkanes deposited within these two hotspots are biodegraded. In contrast, a considerable amount of alkanes still remains near the middle ring of the chamber. Similar tempo-spatial 265patterns are observed for PAHs as well but at much lower biodegradation rate (Figure 5B). 266

Compared to alkanes, biodegradation of PAHs is negligible within the first 100 days. After 267 300 days, biodegradation of PAHs is around 15% near its hotspots, and negligible at other 268 locations. Oil biodegradation greatly alters benthic oxygen conditions (Figure 5C). After 30 269 270 days, the near-surface oxygen level significantly drops within the exfiltration zone to a 271 concentration less than 1.0 mg/L, resulting in anoxic conditions for oil biodegradation. The 272 oxygen level beneath the oiled sediment layer gradually decreases with time, due to large 273 oxygen consumption from the upper oiled layer. In contrast, the oxygen content within the infiltration zone always remains above 2.0 mg/L, which provides a favorable aerobic 274 275condition for oil biodegradation. This is most likely because direct recharge of oxygen-rich seawater to some extent compensates for the oxygen consumption there. Compared to oxygen, 276 277 the nutrient (nitrogen) condition remains at its original level. This is probably due to 278 recycling of nutrients during the oil biodegradation processes (Figure 5D). On the one hand, 279 oil degraders consume nitrogen for microbial growth, but on the other hand, they release nitrogen during their endogenous decay. 280

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Figure 6A reports the alkane biodegradation at three different radial locations. As expected, 282 283 the highest biodegradation rate is observed at the outer ring of the chamber (i.e., R = 45 cm), where above 98% of alkanes are biodegraded after 300 days. The biodegradation rate tends to 284285 be lower as it moves to the inner side (i.e., R = 15 cm), where alkanes are biodegraded about 286 80% after 300 days. The lowest alkane biodegradation occurs near the mid-ring location (i.e., R = 30 cm); nearly 10% of alkanes are biodegraded within the first 20 days, and then the 287 biodegradation rate dramatically drops. Similar tempo-spatial patterns are also observed for 288 289 PAHs at a much lower rate. This is consistent with tempo-spatial variation of oxygen level within the benthic sediment (Figure 6C). The oxygen concentration demonstrates a sudden 290 291 decrease after the oil deposition. In particular, the oxygen concentration drops below 2 mg/L at locations R = 15 cm and 30 cm, indicating anoxic environments there for oil biodegradation. In contrast, oxygen concentration at location R = 45 cm only drops down to 4.5 mg/L and then recovers back to 5.5 mg/L after 300 days.

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Figure 7 reports the percentage (%) loss of alkanes and PAHs at different radial locations. 296 297 Obviously, the % loss is the largest near the outer ring of the chamber (i.e., between R = 40cm and 50 cm), which is about 100% and 20% for alkanes and PAHs, respectively. 298 Comparable loss of alkanes and PAHs is found at the zone between R = 5 cm and 15 cm. 299 300 The % loss drops as it moves to the middle ring of the chamber, reaching the minimum at R = 30~35 cm, which is about 35% and 5% for alkanes and PAHs, respectively. The minimum 301 302 loss is also observed at the center of the chamber (i.e., between R = 0 and 5 cm), which is 303 probably due to persistent low oxygen content there.

304

Pore-water advection significantly affects oil biodegradation in benthic sediments. The pore-305 306 water velocity contour shows a higher magnitude near the surface at the outer and inner ring of the chamber (Figure 8A). Interestingly, it is inconsistent with the magnitude of pressure 307 308 imposed at the sediment surface, which gradually decreases along the radial direction from the outer ring boundary to the inner center of the chamber. This is due to the fact that the 309 310 intensity of pore-water flow is determined by the pressure gradient rather than pressure itself. 311 Figures 8B and 8C show exchange flux and flow across the sediment surface, respectively. Seawater infiltration occurs near the outer ring of the chamber between R = 40 cm and 50 cm, 312 showing the largest rate at the chamber edge. In contrast, groundwater discharge occurs at the 313 314 inner part of the chamber between R = 0 to 40 cm, reaching the highest rate at R = 25 cm. Although the majority of groundwater discharge appears near the middle ring of the chamber, 315discharge flux tends to be the largest near the inner center of the chamber (i.e., R = 5 cm), 316

and gradually decreases as the radius increases. It indicates a relatively poor hydraulic
exchange at the middle ring of the chamber, which to some extent explains the relatively low
oil biodegradation rate there.

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321 The interactions between benthic solute exchange and biodegradation of deposited oil in 322 benthic sediments are dynamic. Figure 9A shows the temporospatial evolution of oxygen 323 exchange across the water-sediment interface after the oil deposition. Although there is a steady oxygen recharge into the benthic sediments, the oxygen exchange curves tend to be 324 325 flat at 0.0 mg/h along the entire discharge zone following the oil deposition (e.g., at time t =10 days and 100 days). It indicates that oil deposition impedes discharge of oxygen from the 326 benthic sediments due to its large consumption for hydrocarbon biodegradation. The positive 327 328 discharge of oxygen starts to occur again after 300 days of the oil deposition at locations 329 between R = 30 cm and 40 cm which is in proximity of the oxygen recharge zone. It indicates 330 that the oxygen concentration is returned to its initial levels by seawater infiltration after 331 major amounts of oil are removed by biodegradation along the discharge flow paths. Figure 332 9B shows the temporospatial evolution of oxygen flux at the bottom of the oiled layer (i.e., Z = 95 cm). The oxygen replenishment along the entire bottom of the oiled sediment layer 333 334 largely decreases within the first 100 days following the deposition, particularly next to the infiltration zone between R = 30 cm and 40 cm. This is due to the oxygen consumption at the 335 336 infiltration zone, diminishing the replacement of oxygen downstream. Such oxygen depletion has lesser impacts on farther discharge locations (e.g., between R = 5 cm and R = 30 cm) due 337 to the following reasons: 1) initial oxygen storage within benthic sediments provides 338 additional oxygen replenishment for downstream oil biodegradation, and 2) the higher water 339 340 and oxygen flux at the farther discharge locations. The intensity of oxygen replenishment is 341 consistent with the magnitude of oil biodegradation within the benthic sediments, indicating the importance of solute (i.e., oxygen) exchange in biodegradation of deposited oil in benthic
sediments.

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### 345 **4. Discussion**

Our results reveal a significant role of benthic flow advection and solute exchange in the fate 346 of oil deposited within permeable marine sediments. Pore-water advection has been 347 recognized as one of the most important drivers for benthic biogeochemical processes 348 [Janssen et al., 2005b; Cook et al., 2007; Rocha, 2008]. Our simulation results indicate that 349 bottom current flows deflected by topographical sediment structures will create subsurface 350 351 pore-water recirculation cells whereby seawater infiltrates into sediment ripple troughs while 352 groundwater discharges near the crest. It leads to a dynamic hydraulic and biogeochemical 353 environment for hydrocarbon biodegradation in benthic sediments. At sediment troughs, the 354 determinants of oil biodegradation rate are infiltration rate and biogeochemical properties of seawater, while biodegradation of oil deposited near the sediment crest is primarily controlled 355 by upwelling groundwater flux beneath the exfiltration zone. Therefore, oil deposited within 356 marine sediments demonstrates spatially different biodegradation rates. Faster oil 357 358 biodegradation occurs in sediment ripple troughs due to direct and sufficient oxygen recharge 359 by seawater infiltration. In contrast, biodegradation of oil deposited uphill is slow and even 360 anoxic (i.e., extremely slow). Such spatial variations will increase the level of challenge to assess oil spill contamination in marine sediments solely based on field measurements. This 361 362 is because random selection of oil-contaminated sediment samples might cause large variability of oil measurements as natural attenuation of oil behaves differently in marine 363 sediments, depending on local topographic and flow patterns. Our results suggest that 364 365 characterization of sediment topographic properties and hydraulic conditions at sampling locations are essential for thorough data interpretation and assessment of oil spills 366

contaminating marine sediments.

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Dissolved oxygen and nutrients are key factors affecting hydrocarbon biodegradation; limited 369 availability of either could slow down the biodegradation rate considerably [Passow and 370 371 Overton, 2021; Wang et al., 2021]. It has been observed that dissolved oxygen concentration 372 affects aerobic microbial growth in a sharp way [Borden and Bedient, 1986; Borden et al., 1986; Chiang et al., 1989]. The optimal oxygen concentration for microbial activities to 373 support aerobic oil biodegradation is 2.0-3.0 mg/L, above which increasing the oxygen 374 375 concentration does not enhance the microbial activities. However, when the concentration drops below 2.0 mg/L, microbial activity switches sharply from aerobic to anaerobic in a way 376 377 that dramatically decreases oil biodegradation rate. Seawater infiltration into marine 378 sediments has been found to provide a major source of dissolved oxygen that enhances 379 sedimentary respiration as well as hydrocarbon biodegradation [Anschutz et al., 2009; Santos 380 et al., 2009]. Our results demonstrate that pore-water advection provides a critical control on 381 benthic solute (e.g., oxygen and nutrients) exchange. Ripple-flow interactions generate an 382 oxygen-rich plume beneath the benthic sediment surface, which has strong implications on 383 the fate and biodegradation of oil deposited in marine sediments. It likely creates an oxygenrich environment (i.e., above 2 mg/L) at ripple troughs that is favorable for aerobic oil 384 385 biodegradation. In contrast, oil biodegradation becomes slow and even anoxic near the ripple 386 crests because of the fact that oxygen is primarily consumed by oil biodegradation near the recharge zone (i.e., ripple troughs) which reduces downstream oxygen replenishment. Our 387 results also show that oil deposition restructures benthic oxygen conditions. Oil 388 389 biodegradation reduces subsurface oxygen content, and consequently impedes discharge of oxygen from the water-sediment interface. Such change could have long-term impacts on 390 benthic microbial community structures which have a large potential to alter nitrogen 391

dynamics, especially inorganic nitrogen release and denitrification processes [*Giblin et al.*,
1995; *Werner et al.*, 2006].

394

### 395 **5. Conclusion**

396 In this paper, we conducted numerical simulations of flow and solute transport processes and hydrocarbon biodegradation in a cylindrical benthic chamber system to investigate influences 397 398 of benthic flow and solute exchange on oil biodegradation in marine sediments. Our results show that ripple-flow interactions create subsurface recirculation cells whereby seawater 399 400 infiltrates into the benthic sediments through ripple troughs while groundwater discharges near ripple crests. It leads to spatially varied oil biodegradation rate in marine sediments, 401 depending on local topographic and flow patterns. Significant oil biodegradation will occur in 402 403 sediment ripple troughs due to direct and sufficient oxygen recharge by seawater infiltration. 404 In contrast, biodegradation of oil deposited uphill is slow and under anoxic condition where oxygen replenishment is prohibited. Overall, this study reveals endogenous mechanisms 405 406 within permeable benthic sediments that affect persistence and biodegradation of the deposited oil. It highlights the importance of benthic flow advection and solute exchange on 407 characterizing fate of oil after its deposition within marine sediments. In particular, benthic 408 flow recirculation along with solute exchange (e.g., oxygen and nutrients) creates a highly 409 410 dynamic environment that increases spatial variability of oil decomposition within marine 411 sediments. In coastal environments, permeability of sediments is heterogeneous, which could strongly alter magnitude and pathways of pore-water flows. In addition, relatively low 412 temperature and high pressure in deep ocean could play an important role in benthic oil 413 414 biodegradation there. These factors are not considered in current work, which are essential and need to be included in future studies. 415

416

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- 421

### 422 Figure Captions

Figure 1. Schematic of oil deposition and benthic exchange processes within permeable marine sediments. Note that the pressure gradients generated by ocean currents over ripples create circulation cells in marine sediments with flows infiltrating in the ripple troughs and exfiltrating from crests.

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428 Figure 2. (A) a comparison of normalized radioactivity strength measured by PET (gray-429 scale filled contours) in Khalili et al., (1999) with numerical results obtained from Basu and Khalili (1999) and CYL-BIOMARUN model at different times. The dashed and solid curves 430 represent the simulated radioactivity of 0.5 and 0.9, respectively. (B) A comparison of 431 washout depths obtained from experiments and numerical simulations. Note that in order to 432 compensate for the effect of premixing (i.e., observed loss of some dye from the sediment to 433 the fluid layer at the initial instant of the experiments) in the experiments, Khalili et al., (1999) 434 shifted back their numerical results by 12 minutes for additional comparison, plotted as a 435 dashed line. 436

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Figure 3. Simulated steady-state distributions of (A) pressure, (C) dissolved oxygen, and (D) nutrient (i.e., nitrogen) in the benthic chamber system. (B) Initial oil (i.e., alkanes and PAHs) distribution in the benthic sediment. In Figure 3A, the upper panel shows the pressure distribution along the water-sediment interface, and the lower panel shows steady-state pressure increment (above 100 cm) built by pressure gradients imposed at the water-sediment interface.

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Figure 4. (A) Simulated flow paths of the particles released at 0.1-m interval along the sediment surface from R = 2 cm to 50 cm. (B) Travel time of the particles. Note that backward particle tracking was used to track advective flow paths. The start and end locations of particle flow paths are marked with blue symbols and a red line, respectively.

Figure 5. Simulated concentration contours of (A) alkanes, (B) PAHs, (C) oxygen, and (D) nutrient at different times. The dashed contour lines for alkanes and PAHs represent 75% and 95% of its initial concentration, respectively. The threshold of 2.0 mg/L that supports aerobic oil biodegradation is marked in oxygen contours. The nutrient contours demonstrate a negligible change with time; therefore, only its 300-day concentration contour is shown herein.

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Figure 6. Temporal change of concentration for (A) alkanes, (B) PAHs, and (C) dissolved oxygen at three different radial locations (Z = 97.5 cm, and R = 15 cm, 30 cm, and 45 cm). The output locations are marked in the upper panel using the same color as the concentration curves.

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- 463 **Figure 7.** Percentage loss of (A) alkanes and (B) PAHs at different radial locations.
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Figure 8. (A) Simulated pore-water velocity field. The spatial distribution of (B) exchange flux (cm<sup>3</sup>/h·cm<sup>2</sup>) and (C) exchange flow (cm<sup>3</sup>/h) along the sediment-water interface. In Figure 8A, the color contour and uniform vectors represent magnitude and direction of pore water flow, respectively.

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Figure 9. (A) Oxygen exchange across the water-sediment interface. (B) Oxygen flux across bottom of the oiled layer (i.e., Z = 95 cm). Note that the positive and negative values represent upward and downward flow directions, respectively.

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